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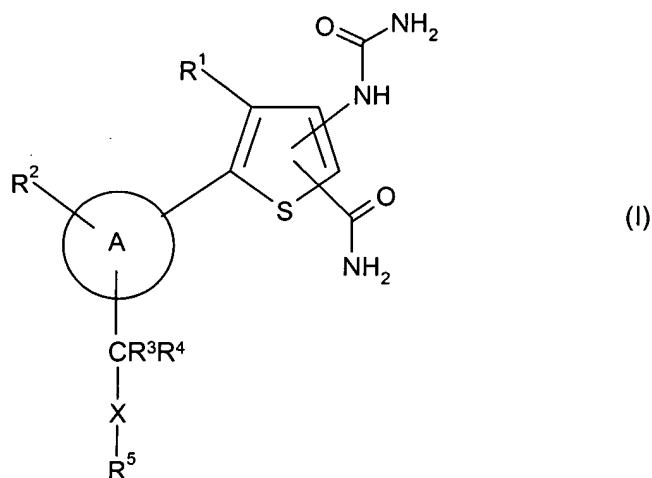
JC17 Rec'd PC17/10 13 JUL 2005

Amendments to the Claims:

This listing of claims replaces all prior versions and listings of claims in the application:

Listing of Claims:

1. (Original) A compound of formula (I)



wherein

$\text{R}^1$  represents H or  $\text{CH}_3$ ;

$\text{R}^2$  represents hydrogen, halogen, cyano, C1 to 2 alkyl, trifluoromethyl or C1 to 2 alkoxy;

$\text{R}^3$  and  $\text{R}^4$  independently represent H or  $\text{CH}_3$ ;

or the group  $\text{CR}^3\text{R}^4$  together represents a C3 to 6 cycloalkyl ring;

$\text{A}$  represents a six-membered aromatic ring optionally incorporating one or two nitrogen atoms; and the group  $-\text{CR}^3\text{R}^4-\text{X}-\text{R}^5$  is bonded to ring  $\text{A}$  in the 4-position relative to the thiophene ring;

$\text{X}$  represents  $\text{NR}^6$ ;

R<sup>5</sup> represents H, C1 to 6 alkyl, C2 to 6 alkenyl or C3 to 6 cycloalkyl; said cycloalkyl group optionally incorporating one heteroatom selected from O, S(O)<sub>n</sub> or NR<sup>7</sup>; said alkyl group being optionally further substituted by one or more groups selected independently from CN, OH, C1 to 4 alkoxy, F, a C5 to 10 monocyclic or bicyclic aromatic ring system optionally incorporating one or two heteroatoms independently selected from O, S and N, and said ring system being optionally further substituted by one or more substituents selected independently from halogen, C1 to 2 alkyl, C1 to 2 alkoxy or CF<sub>3</sub>; or said alkyl being optionally further substituted by a C5 to 6 cycloalkyl ring that optionally incorporates a heteroatom selected from O, S(O)<sub>m</sub> or NR<sup>8</sup> and/or a carbonyl group and is optionally further substituted by OH;

R<sup>6</sup> represents H or C1 to 6 alkyl; said alkyl group being optionally further substituted by CN, OH, C1 to 4 alkoxy or one or more fluoro atoms;

n and m independently represent an integer 0, 1 or 2;

R<sup>7</sup> and R<sup>8</sup> independently represent H or C1 to 2 alkyl;

and pharmaceutically acceptable salts thereof.

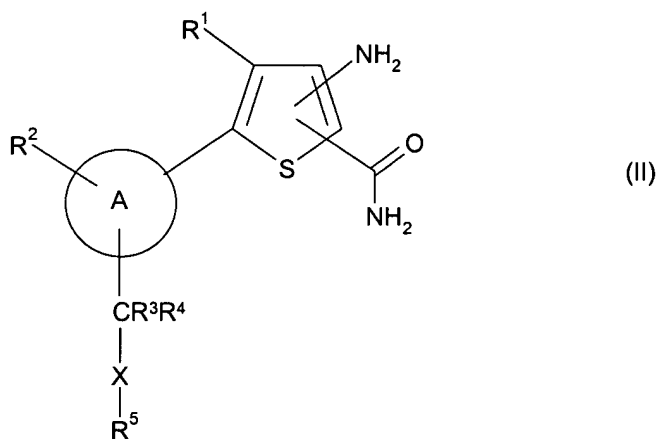
2. (Original) A compound of formula (I), according to Claim 1, wherein R<sup>1</sup> represents H.

3. (Currently amended) A compound of formula (I), according to Claim 1 ~~or Claim 2~~, in which A represents optionally substituted phenyl.

4. (Currently amended) A compound of formula (I), according to ~~any one of Claims 1 to 3~~ Claim 1, in which R<sup>3</sup> and R<sup>4</sup> each represent H.

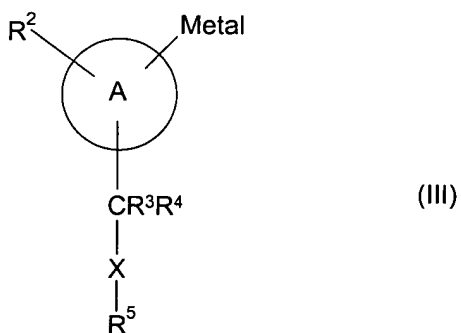
5. (Currently amended) A process for the preparation of a compound of formula (I), according to ~~any one of Claims 1 to 4~~ Claim 1, which comprises:

(a) reaction of a compound of formula (II):

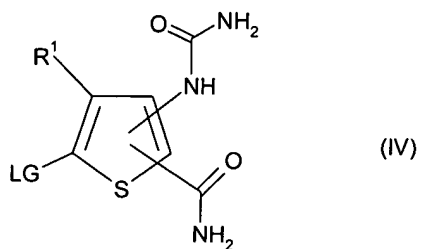


wherein A, R<sup>1</sup>, R<sup>2</sup>, R<sup>3</sup>, R<sup>4</sup>, R<sup>5</sup> and X are as defined in Claim 1 with an isocyanate; or

(b) reaction of a compound of formula (III)

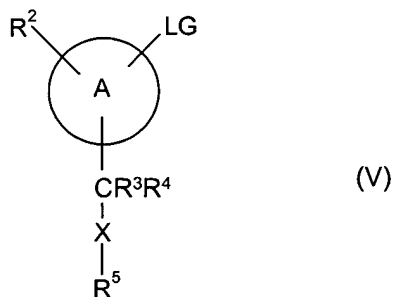


wherein A, R<sup>2</sup>, R<sup>3</sup>, R<sup>4</sup>, R<sup>5</sup> and X are as defined in Claim 1,  
 with a compound of formula (IV)



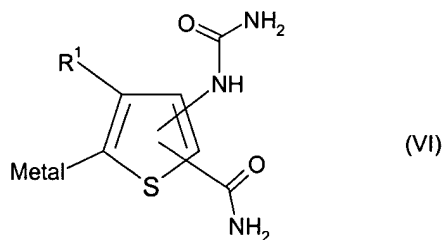
wherein R<sup>1</sup> is as defined in Claim 1 and LG represents a leaving group; or

(c) reaction of a compound of formula (V)



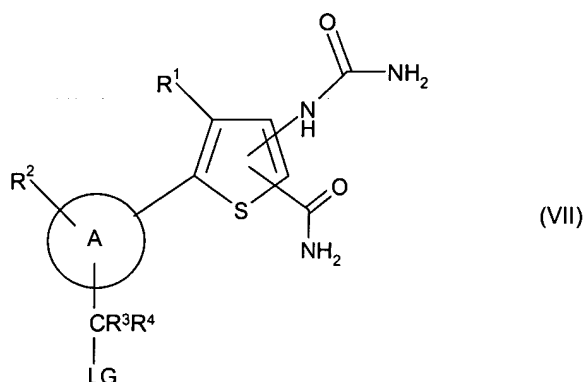
wherein A,  $R^2$ ,  $R^3$ ,  $R^4$ ,  $R^5$  and X are as defined in Claim 1 and LG represents a leaving group,

with a compound of formula (VI)



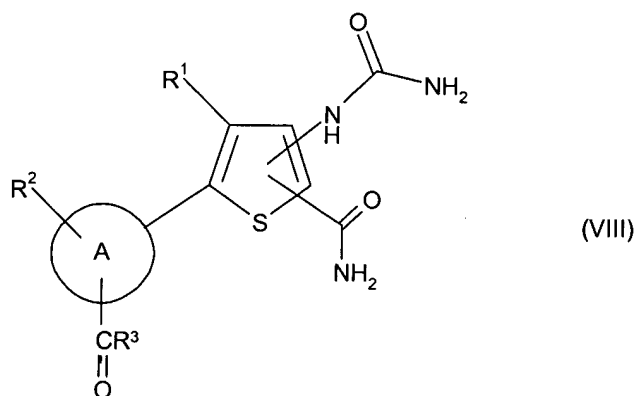
wherein  $R^1$  is as defined in Claim 1; or

(d) reaction of a compound of formula (VII)

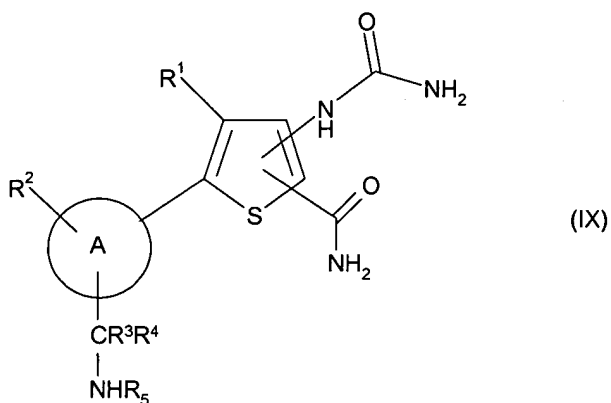


wherein A, R<sup>1</sup>, R<sup>2</sup>, R<sup>3</sup> and R<sup>4</sup> are as defined in Claim 1, and LG represents a leaving group,

with an amine of formula R<sup>5</sup>R<sup>6</sup>NH, wherein R<sup>5</sup> and R<sup>6</sup> are as defined in Claim 1; or  
 (e) reaction of a compound of formula (VIII)



wherein A, R<sup>1</sup>, R<sup>2</sup> and R<sup>3</sup> are as defined in Claim 1,  
 with an amine of formula R<sup>5</sup>R<sup>6</sup>NH wherein R<sup>5</sup> and R<sup>6</sup> are as defined in Claim 1, under reductive amination conditions; or  
 (f) reaction of a compound of formula (IX)



wherein R<sup>1</sup>, R<sup>2</sup>, R<sup>3</sup>, R<sup>4</sup>, R<sup>5</sup> and A are as defined in Claim 1,  
with an aldehyde or ketone under reductive amination conditions;  
and where necessary converting the resultant compound of formula (I), or another salt thereof, into a pharmaceutically acceptable salt thereof; or converting the resultant compound of formula (I) into a further compound of formula (I); and where desired converting the resultant compound of formula (I) into an optical isomer thereof.

6. (Currently amended) A pharmaceutical composition comprising a compound of formula (I), or a pharmaceutically acceptable salt thereof, as claimed in ~~any one of claims 1 to 4~~ Claim 1 in association with a pharmaceutically acceptable adjuvant, diluent or carrier.

7. (Currently amended) A pharmaceutical composition adapted for administration by inhalation or insufflation[[.]] comprising a compound of formula (I), or a pharmaceutically acceptable salt thereof, as claimed in ~~any one of claims 1 to 6~~ Claim 1 in association with a pharmaceutically acceptable adjuvant, diluent or carrier.

8. (Currently amended) A process for the preparation of a pharmaceutical composition ~~as claimed in Claim 6~~ which comprises mixing a compound of formula (I), or a pharmaceutically acceptable salt thereof, as claimed in ~~any one of claims 1 to 4~~ Claim 1 with a pharmaceutically acceptable adjuvant, diluent or carrier.

9. (Cancelled)

10. (Cancelled)

11. (Currently amended) A method for the treatment or prophylaxis of inflammatory disease, comprising administering to a person suffering from or at risk of said disease a therapeutically effective amount [[Use]] of a compound of formula (I), or a pharmaceutically acceptable salt thereof, as claimed in Claim 1 ~~any one of claims 1 to 4 in the manufacture of a medicament for use in the treatment or prophylaxis of inflammatory disease.~~

12. (Currently amended) The [[use]] method as claimed in Claim 11 wherein the disease is rheumatoid arthritis.

13. (Currently amended) The [[use]] method as claimed in Claim 11 wherein the disease is chronic obstructive pulmonary disease.

14. (Cancelled)

15. (Currently amended) A method of treating, or reducing the risk of, ~~diseases a~~ disease or conditions condition in which inhibition of IKK-2 activity is beneficial which comprises administering to a person suffering from or at risk of said disease or condition a therapeutically effective amount of a compound of formula (I), or a pharmaceutically acceptable salt thereof, as claimed in ~~any one of claims 1 to 4~~ Claim 1.

16. (New) A compound of formula (I), according to Claim 2, in which A represents optionally substituted phenyl.

17. (New) A compound of formula (I), according to Claim 2, in which  $R^3$  and  $R^4$  each represent H.

18. (New) A compound of formula (I), according to Claim 3, in which  $R^3$  and  $R^4$  each represent H.

19. (New) A compound of formula (I), according to Claim 16, in which  $R^3$  and  $R^4$  each represent H.

20. (New) The method as claimed in Claim 15 wherein the disease is cancer.